RECEIVED CENTRAL FAX CENTER

IN THE CLAIMS

1. (currently amended) A compound of formula 1

$$R^3$$
 R^4
 $N^{\pm 0}$
 R^5
 R^5
 R^1
 R^1
 R^2
 R^4
 R^5
 R^5

wherein

R1

(i) is -C1-10-alkyl, straight-chain or branched-chain, optionally mono- or polysubstituted by -OH, -SH,

 $-NH_2$, $-NHC_{1-6}$ -alkyl, $-N(C_{1-6}$ -alkyl)₂, $-NHC_{6-14}$ -aryl,

 $-N(C_{6-14}-aryl)_2$, $-N(C_{1-6}-alkyl)(C_{6-14}-aryl)$, $-NO_2$,

-CN, -F, -Cl, -Br, -I, -O-C₁₋₆-alkyl, -O-C₆₋₁₄-aryl, -S-C₁₋₆-alkyl, -S-C₆₋₁₄-aryl, -SO₃H, -SO₂C₁₋₆-alkyl,

 $-SO_2C_{6-14}$ -aryl, $-OSO_2C_{1-6}$ -alkyl, $-OSO_2C_{6-14}$ -aryl,

-COOH, -(CO)C₁₋₅-alkyl, -COO-C₁₋₅-alkyl, -O(CO)C₁₋₅-alkyl, by a mono-, bi- or tricyclic saturated or mono- or polyunsaturated carbocycle earbocycles with 3-14 ring members or/and by mono, bi or tricyclic saturated or mono or polyunsaturated heterocycles with 5-15 ring members and 1-6 heteroatoms, which are preferably N, O and S,

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where the C₆₋₁₄-aryl groups and the carbocyclic and beterocyclic substituents in turn may optionally be substituted at least once one or more times by -C₁₋₆-alkyl, -OH, -NH₂, -NHC₁₋₆-alkyl, -N(C₁₋₆-alkyl)₂, -NO₂, -CN, -F, -Cl, -Br, -I, -O-C₁₋₆-alkyl, -S-C₁₋₆-alkyl, -SO₂C₁₋₆-alkyl, -OSO₂C₁₋₆-alkyl, -COOH, -(CO)C₁₋₅-alkyl, -COO-C₁₋₅-alkyl or or/and -O(CO)C₁₋₅-alkyl, and wherein where the alkyl groups on the carbocyclic and heterocyclic substituents in turn may optionally be substituted one-or more times at least once by -OH, -SH, -NH₂, -F, -Cl, -Br, -I, -SO₃H or or/and -COOH, or

(ii) is $-C_{2-10}$ -alkenyl, mono- or polyunsaturated, straight-chain or branched-chain, optionally mono- or polysubstituted by <u>at least one of</u> -OH, -SH, -NH₂, -NHC₁₋₆-alkyl, -N(C₁₋₆-alkyl)₂, -NHC₆₋₁₄-aryl, -N(C₆₋₁₄-aryl)₂, -N(C₁₋₆-alkyl)(C₆₋₁₄-aryl), -NO₂, -CN, -F, -Cl, -Br, -I, -O-C₁₋₆-alkyl, -O-C₆₋₁₄-aryl, -S-C₁₋₆-alkyl, -S-C₆₋₁-aryl, -SO₂H, -SO₂C₁₋₆-alkyl, -SO₂C₁₋₆-alkyl, -OSO₂C₆₋₁₄-aryl, -COOH, -(CO)C₁₋₅-alkyl, -COO-C₁₋₅-alkyl <u>or or/and</u> -O(CO)C₁₋-alkyl, by mono-, bi- or tricyclic saturated or mono- or <u>a</u> polyunsaturated <u>carbocycle</u> <u>earbocycles</u> with 3-14 ring members <u>or/and</u> by mono-, bi- or tricyclic saturated or mono- or polyunsaturated heterocycles with 5-15 ring members and 1-6 heteroctoms, which are preferably N, O and S,

where the C_{6-14} -aryl groups and the carbocyclic and heterocyclic substituents in turn may optionally be substituted at least once one or more times by $-C_{1-6}$ -alkyl, -OH, $-NH_2$, $-NHC_{1-6}$ -alkyl, $-N(C_{1-6}$ -alkyl)₂, $-NO_2$, -CN, -F, -Cl, -Br, -I, $-O-C_{1-6}$ -alkyl, $-S-C_{1-6}$ -alkyl, $-SO_2C_{1-6}$ -alkyl, $-OSO_2C_{1-6}$ -alkyl, -COOH, $-(CO)C_{1-5}$ -alkyl, $-COO-C_{1-5}$ -alkyl, $-OCO-C_{1-5}$ -alkyl, $-OCO-C_{1-5}$ -alkyl,

and where the alkyl groups on the carbocyclic and heterocylic substituents in turn may optionally be substituted at least once one or more times by -OH, -SH, -NH₂, -F, -Cl, -Br, -I, -SO₃H or or/and -COOH,

R² is hydrogen or -C₁₋₃-alkyl,

R³ is a hydroxyl group, and wherein

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R⁴ and R⁵ may be identical or different and are hydrogen, -C₁₋₆-alkyl, -OH, -SH, -NH₂, -NHC₁₋₆-alkyl, -N(C₁₋₆-alkyl)₂, -NO₂, -CN, -SO₂H, -SO₃-C₁₋₆-alkyl, -COOH, -COO-C₁₋₆-alkyl, -O(CO)-C₁₋₅-alkyl, -F, -Cl, -Br, -I, -O-C₁₋₆-alkyl, -S-C₁₋₆-alkyl, or -phenyl erpyridyl, where the phenyl substituent or pyridyl substituents in turn may optionally be substituted at least once one or more times by -C₁₋₃-alkyl, -OH, -SH, -NH₂, -NHC₁₋₃-alkyl, -N(C₁₋₃-alkyl)₂, -NO₂, -CN, -SO₃H, -SO₃C₁₋₃-alkyl, -COOH, -COOC₁₋₃-alkyl, -F, -Cl, -Br, -I, -O-C₁₋₃-alkyl, -S-C₁₋₃-alkyl, or or/and—O(CO)C₁₋₃-alkyl, and where the alkyl substituents in turn may optionally be substituted at least once one or more times by -OH, -SH, -NH₂, -F, -Cl, -Br, -I, -SO₃H, -SO₃C₁₋₃-alkyl, -COOH, -COOC₁₋₃-alkyl, -O-C₁₋₃-alkyl, -S-C₁₋₃-alkyl or or/and -O(CO)-C₁₋₃-alkyl, or a salt thereof theteof.

- 2. (previously presented) A compound as claimed in claim 1, having at least one asymmetric carbon atom in the D form, the L form and D,L mixtures, and in the case of a plurality of asymmetric carbon atoms also the diaster-comeric forms.
- 3. (previously presented) A compound as claimed in claim 1, wherein R² is hydrogen or a methyl group.
 - 4. (canceled)
- 5. (currently amended) A compound as claimed in claim 1, selected from the group consisting of:

N-(3,5-dichloro-1-oxopyridiu-4-yl)-[1-(4-fluorobenzyl)-5-hydroxyindol-3-yl]glyoxylamide;

N-(3,5-dichloro-1-oxopyridin-4-yl)-[1-(4-chlorobenzyl)-5-hydroxyindol-3-yl]glyoxylamide;

N-(1-oxopyridin-4-yl)-[1-(4-fluorobenzyl)-5-hydroxyindol-3-yl]glyoxylamide;

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N-(3,5-dichloro-1-oxopyridin-4-yl)-[1-(2,4-dichlorobenzyl)-5-hydroxyindol-3yl]glyoxylamide;

N-(3,5-dichloro-1-oxopyridin-4-yl)-[5-hydroxy-1-(3-nitrobenzyl)-indol-3yl]glyoxylamide;

N-(3,5-dichloro-1-oxopyridin-4-yl)-[1-(2,6-difluorobenzyl)-5-hydroxyindol-3yl]glyoxylamide;

N-(3,5-dichloro-1-oxopyridin-4-yl)-(5-hydroxy-1-isobutylindol-3yl)glyoxylamide;

N-(3,5-dichloro-1-oxopyridin-4-yl)-(1-cyclopropyl-methyl-5-hydroxyindol-3yl)glyoxylamide;

N-(3,5-dichloro-1-oxopyridin-4-yl)-[5-hydroxy-1-(4-hydroxybenzyl)-indol-3yl]glyoxylamide;

N-(3,5-dichloro-1-oxopyridin-4-yl)-N-methyl-[1-(4-fluorobenzyl)-5hydroxyindol-3-yl]glyoxylamide;

and or a physiologically tolerated salt salts thereof.

6. (previously presented) The compound of claim 1 that is N-(3,5-dichloro-1oxopyridin-4-yl)-[1-(4-fluorobenzyl)-5-hydroxyindol-3-yl]glyoxylamide.

Claims 7-15 (canceled)

16 (previously presented) A compound as claimed in claim 2 wherein R² is hydrogen or a methyl group.

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17.(previously presented) The compound of claim 1 that is a physiologically acceptable salt of N-(3,5-dichloro-1-oxopyridin-4-yl)-[1-(4-fluorobenzyl)-5-hydroxyindol-3-yl]glyoxylamide.

Claims 18-19 (canceled)

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20.(currently amended) A compound as claimed in claim 1, wherein at least one of R^4 and R^5 is F, Cl, Br, or I a hologen-atom.

21. (new) A pharmaceutical composition comprising the compound as claimed in claim 1 and at least one of a conventional physiologically tolerated carrier, diluent or excipient.